```
> d his

     (FILE 'HOME' ENTERED AT 10:51:57 ON 03 JUN 2004)

     FILE 'CAPLUS' ENTERED AT 10:53:58 ON 03 JUN 2004
L1             0 S WO20059486/PN
L2             1 S WO200059486/PN
                 SELECT L2 1 RN
L3        110742 S E1-E81
L4          1188 S L3 AND ?PROPANOL

     FILE 'REGISTRY' ENTERED AT 10:57:36 ON 03 JUN 2004
L5             1 S 136817-59-9/RN
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                 SET NOTICE LOGIN DISPLAY

     FILE 'REGISTRY' ENTERED AT 10:58:59 ON 03 JUN 2004
L6             1 S 134234-12-1/RN
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     FILE 'REGISTRY' ENTERED AT 11:03:11 ON 03 JUN 2004
L7             1 S 133454-47-4/RN
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     FILE 'CAPLUS' ENTERED AT 11:03:24 ON 03 JUN 2004
L8            46 S L6
L9             2 S L8 AND NK1
L10           51 S L6 OR CP(W)(101606 OR 98113) OR TRAXOPRODIL
L11           15 S L10 NOT PY>=1999
                 S 145742-28-5/REG# OR  QUINIDINE OR CHINIDIN OR CONCHININ? OR C

     FILE 'REGISTRY' ENTERED AT 11:25:08 ON 03 JUN 2004
L12            1 S 18786-24-8/RN

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L13          251 S L12

     FILE 'REGISTRY' ENTERED AT 11:25:09 ON 03 JUN 2004
L14            1 S 145742-28-5/RN

     FILE 'CAPLUS' ENTERED AT 11:25:09 ON 03 JUN 2004
L15           58 S L14
L16         7045 S L15 OR QUINIDINE OR CHINIDIN OR CONCHININ? OR CONQUININE OR C
L17            2 S L10 AND L16
L18         2342 S SERTRALINE OR VENLAFAXINE OR DEXMEDETOMIDINE OR TRIPENNELAMIN
L19            1 S L18 AND L10
L20          549 S L18 AND NEURO?
L21          217 S L20 NOT PY>=1999
L22            6 S L21 AND US/PC

     FILE 'REGISTRY' ENTERED AT 11:51:53 ON 03 JUN 2004
                 E  SERTRALINE/CN
L23            1 S E3

     FILE 'CAPLUS' ENTERED AT 11:52:30 ON 03 JUN 2004
L24         1275 S L23 OR SERTRALINE OR CP(W)51974
L25          213 S L24 AND US/PC
L26           43 S L25 AND NEURO?
L27            5 S L26 NOT PY>=1999
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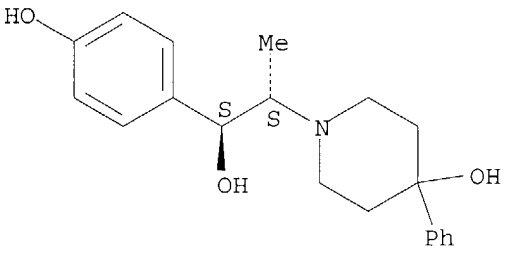
L6 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN
RN **134234-12-1** REGISTRY
CN 1-Piperidineethanol, 4-hydroxy-α-(4-hydroxyphenyl)-β-methyl-4-phenyl-, (αS,βS)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN 1-Piperidineethanol, 4-hydroxy-α-(4-hydroxyphenyl)-β-methyl-4-phenyl-, [S-(R*,R*)]-
OTHER NAMES:
CN (+)-CP 101606
CN CP 101606
CN CP 98113
CN Traxoprodil
FS STEREOSEARCH
MF C20 H25 N O3
CI COM
SR CA
LC STN Files: ADISINSIGHT, BIOSIS, BIOTECHNO, CA, CAPLUS, DDFU, DRUGU, EMBASE, IMSDRUGNEWS, IMSRESEARCH, PHAR, PROMT, PROUSDDR, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
DT.CA CAplus document type: Journal; Patent
RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)
RL.NP Roles from non-patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); USES (Uses)
RLD.NP Roles for non-specific derivatives from non-patents: BIOL (Biological study); USES (Uses)

Absolute stereochemistry. Rotation (+).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

46 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
46 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L2            1 WO200059486/PN
                (WO2000059486/PN)

=> select l2
ENTER ANSWER NUMBER OR RANGE (1-):1
ENTER DISPLAY CODE (TI) OR ?:rn
E1 THROUGH E81 ASSIGNED

=> s e1-e81
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          5538 10540-29-1/BI
           273 106133-20-4/BI
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           538 1131-64-2/BI
           434 113775-47-6/BI
           794 114-86-3/BI
           117 115956-12-2/BI
           157 124937-51-5/BI
           609 125-28-0/BI
           772 125-29-1/BI
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          1409 155213-67-5/BI
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	236	64-13-1/BI
	174	6621-47-2/BI
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	726	68-88-2/BI
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	441	71320-77-9/BI
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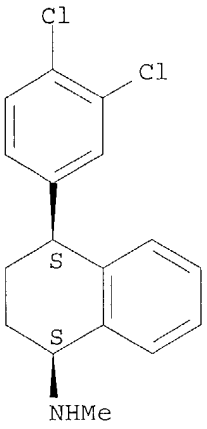
L23 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN
RN 79617-96-2 REGISTRY

Absolute stereochemistry. Rotation (+).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

CN 1-Naphthalenamine, 4-(3,4-dichlorophenyl)-1,2,3,4-tetrahydro-N-methyl-, (1S,4S)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN 1-Naphthalenamine, 4-(3,4-dichlorophenyl)-1,2,3,4-tetrahydro-N-methyl-, (1S-cis)-
OTHER NAMES:
CN (+)-Sertraline
CN CP 51974
CN **Sertraline**

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=> s l10 and l16
L17          2 L10 AND L16

=> d ibib abs 1-2

L17  ANSWER 1 OF 2  CAPLUS  COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:         2000:725447  CAPLUS
DOCUMENT NUMBER:          133:301178
TITLE:                    Use of CYP2D6 inhibitors in combination therapies
INVENTOR(S):              Obach, Ronald Scott
PATENT ASSIGNEE(S):       Pfizer Products Inc., USA
SOURCE:                   PCT Int. Appl., 18 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:            Patent
LANGUAGE:                 English
FAMILY ACC. NUM. COUNT:   1
PATENT INFORMATION:

     PATENT NO.         KIND  DATE           APPLICATION NO.   DATE
     ---------------    ----  --------       ---------------   --------
     WO 2000059486      A2    20001012       WO 2000-IB304     20000320
     WO 2000059486      C1    20020725
         W:  AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
             CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
             IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
             MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
             SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,
             AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     BR 2000009564      A     20020108       BR 2000-9564      20000320
     EP 1242058         A1    20020925       EP 2000-909570    20000320
         R:  AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL
     EE 200100524       A     20021216       EE 2001-524       20000320
     JP 2003523936      T2    20030812       JP 2000-609050    20000320
     US 2003144220      A1    20030731       US 2000-528978    20000321
     HR 2001000722      A1    20020831       HR 2001-722       20011004
     ZA 2001008158      A     20030724       ZA 2001-8158      20011004
     NO 2001004858      A     20011205       NO 2001-4858      20011005
     BG 106075          A     20020628       BG 2001-106075    20011101
     US 2004018253      A1    20040129       US 2003-622301    20030718
     US 2004028755      A1    20040212       US 2003-624123    20030721
PRIORITY APPLN. INFO.:                    US 1999-128136P  P  19990407
                                          WO 2000-IB304    W  20000320
                                          US 2000-528978   A3 20000321
AB   This invention relates to the use of a CYP2D6 inhibitor in combination
     with a drug having CYP2D6-catalyzed metabolism, wherein the drug and the
     CYP2D6 inhibitor are not the same compound; and pharmaceutical compns. for
     said use.

L17  ANSWER 2 OF 2  CAPLUS  COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:         2000:351162  CAPLUS
DOCUMENT NUMBER:          133:790
TITLE:                    New use of glutamate antagonists for the treatment of
                          cancer
INVENTOR(S):              Ikonomidou, Hrissanthi
PATENT ASSIGNEE(S):       Germany
SOURCE:                   Eur. Pat. Appl., 21 pp.
                          CODEN: EPXXDW
DOCUMENT TYPE:            Patent
LANGUAGE:                 English
FAMILY ACC. NUM. COUNT:   1
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PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1002535	A1	20000524	EP 1998-250380	19981028
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
AU 9964750	A1	20000515	AU 1999-64750	19991022
EP 1124553	A1	20010822	EP 1999-952622	19991022
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2002528415	T2	20020903	JP 2000-578005	19991022

PRIORITY APPLN. INFO.: EP 1998-250380 A 19981028
WO 1999-EP8004 W 19991022

AB New therapies can be devised based upon a demonstration of the role of glutamate in the pathogenesis of cancer. Inhibitors of the interaction of glutamate with the AMPA, kainate, or NMDA receptor complexes are likely to be useful in treating cancer and can be formulated as pharmaceutical compns. They can be identified by appropriate screens.

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT